

Mouse monoclonal [SB62a] Anti-Rabbit IgG light chain (HRP) ab99697

★★★★★ [1 Abreviews](#) [78 References](#) [7 图像](#)

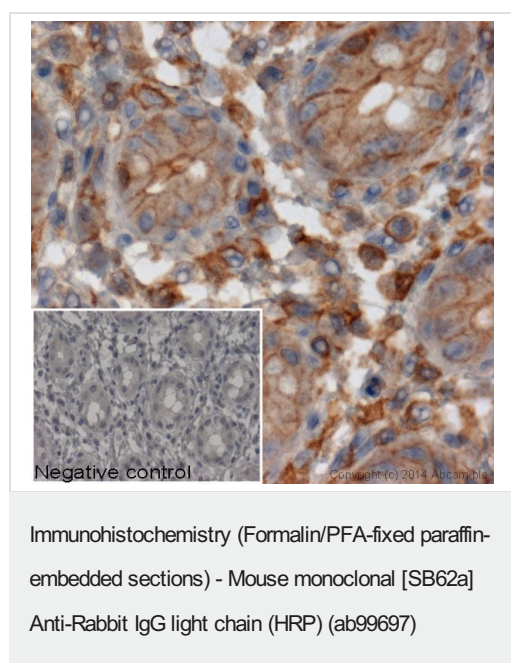
概述	
产品名称	小鼠单克隆抗体[SB62a]抗兔IgG light chain (HRP)
宿主	Mouse
靶标种属	Rabbit
特异性	ab99697 reacts with the light chain of native Rabbit IgG primary antibodies and does not bind to the reduced and denatured Rabbit IgG heavy chain band (50kD).
经测试应用	适用于: IHC-P, ELISA, WB
免疫原	Full length protein corresponding to Rabbit IgG Light chain. Rabbit Light Chain
偶联物	HRP

性能	
形式	Liquid
存放说明	Shipped at 4°C. Store at +4°C.
存储溶液	pH: 7.4 Constituents: 50% Glycerol (glycerin, glycerine), 50% PBS
纯度	Protein A purified
克隆	单克隆
克隆编号	SB62a
同种型	IgG2b
轻链类型	lambda

应用	
The Abpromise guarantee	Abpromise™ 承诺保证使用ab99697于以下的经测试应用
“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。	

应用	Ab评论	说明
IHC-P		1/8000.
ELISA		1/4000 - 1/8000.
WB		Use at an assay dependent concentration. Note: This antibody is not recommended for sensitive or quantitative detection of the reduced and denatured Rabbit Light Chains on Western Blots.

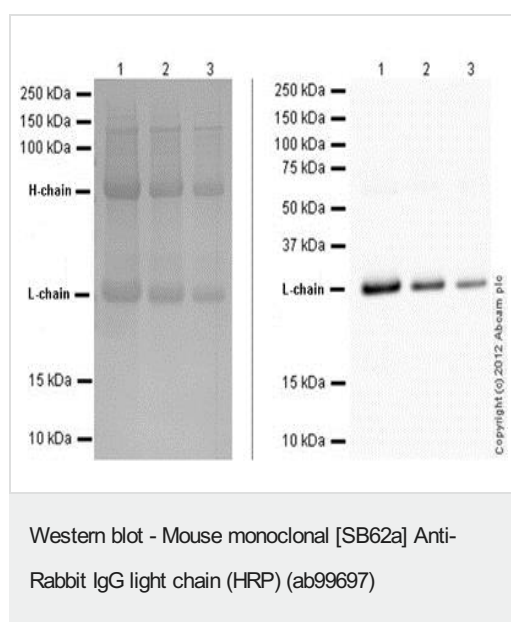
图片



IHC image of beta actin staining in human colon formalin fixed paraffin embedded tissue section*. The section was pre-treated using pressure cooker heat mediated antigen retrieval with sodium citrate buffer (pH6) for 30mins. The section was incubated with **ab8227**, 3µg/ml overnight at +4°C. An HRP-conjugated secondary (ab99697, 1/8000 dilution) was used for 1hr at room temperature. The section was counterstained with haematoxylin and mounted with DPX.

The inset negative control image is secondary-only at 1/4000 dilution.

*Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre



Western blot showing that Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697) specifically binds to light-chains

Left blot: Coomassie stain showing the rabbit heavy (~50kDa) and light-chains (~25 kDa)

Lane 1: 10 µg rabbit monoclonal was loaded

Lane 2: 5 µg rabbit monoclonal was loaded

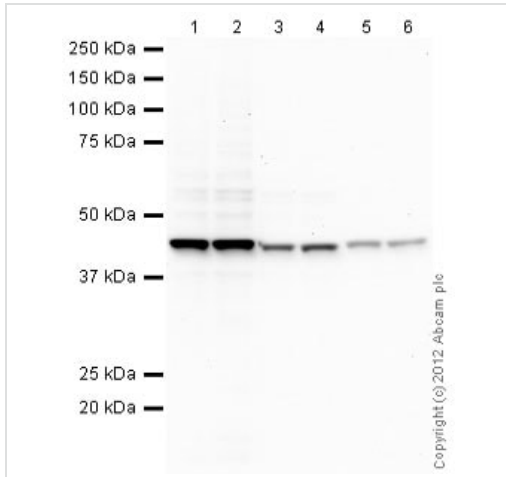
Lane 3: 2 µg rabbit monoclonal was loaded

Right blot: WB using Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697) as secondary antibody at 1/10,000

Lane 1: 1 µg rabbit monoclonal was loaded

Lane 2: 0.5 µg rabbit monoclonal was loaded

Lane 3: 0.25 µg rabbit monoclonal was loaded



Western blot - Mouse monoclonal [SB62a] Anti-Rabbit IgG light chain (HRP) (ab99697)

All lanes : Anti-beta Actin antibody - Loading Control ([ab8227](#)) at 1 µg/ml

Lanes 1-6 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate ([ab27252](#))
Lysates/proteins at 10 µg per lane.

Secondary

Lanes 1 - 2 : Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697) at 1/2000 dilution

Lanes 3 - 4 : Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697) at 1/10000 dilution

Lanes 5 - 6 : Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697) at 1/20000 dilution

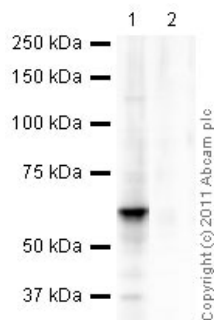
Developed using the ECL technique.

Performed under reducing conditions.



Immunoprecipitation - Mouse monoclonal [SB62a] Anti-Rabbit IgG light chain (HRP) (ab99697)

LAP2 alpha was immunoprecipitated using 0.5mg HeLa whole cell extract, 5µg of Rabbit polyclonal to LAP2 alpha ([ab5162](#)) and 50µl of protein G magnetic beads (lane 1). The antibody was incubated with the Protein G beads for 10min under agitation. No antibody was added to the control (lane 2). HeLa whole cell extract diluted in RIPA buffer was added to each sample and incubated for 10min under agitation. Proteins were eluted by addition of 40µl SDS loading buffer and incubated for 10min at 70°C; 10µl of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with [ab5162](#). Secondary: Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697).
Bands: 83kDa: LAP2 alpha.



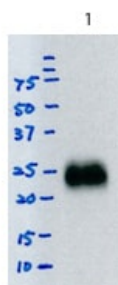
Immunoprecipitation - Mouse monoclonal [SB62a]
Anti-Rabbit IgG light chain (HRP) (ab99697)

Catalase was immunoprecipitated using 0.5mg Hela whole cell extract, 5µg of Rabbit polyclonal to Catalase ([ab1877](#)) and 50µl of protein G magnetic beads (lane 1). The antibody was incubated with the Protein G beads for 10min under agitation. No antibody was added to the control (lane 2). Hela whole cell extract diluted in RIPA buffer was added to each sample and incubated for 10min under agitation. Proteins were eluted by addition of 40µl SDS loading buffer and incubated for 10min at 70°C; 10µl of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with [ab1877](#). Secondary: Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697). Bands: 60kDa: Catalase



Immunoprecipitation - Mouse monoclonal [SB62a]
Anti-Rabbit IgG light chain (HRP) (ab99697)

Rad21 was immunoprecipitated using 0.5mg Hela whole cell extract, 5µg of Rabbit polyclonal to Rad21 and 50µl of protein G magnetic beads (lane 1). The antibody was incubated with the Protein G beads for 10min under agitation. No antibody was added to the control (lane 2). Hela whole cell extract diluted in RIPA buffer was added to each sample and incubated for 10min under agitation. Proteins were eluted by addition of 40µl SDS loading buffer and incubated for 10min at 70°C; 10µl of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with [ab42478](#). Secondary: Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697) .. Band: 100kDa: Rad21; Non specific - 50kDa: We are unsure as to the identity of this extra band. Although the predicted band size is 75kDa based on Swiss-prot data, a band of 120kDa has been previously observed. Mol Cell Biol. 2002 Dec;22(23):8267-77 PMID: 12417729. Th



Western blot - Mouse monoclonal [SB62a] Anti-
Rabbit IgG light chain (HRP) (ab99697)

Lane 1: Rabbit IgG loaded at 1ug/well
ab99697 diluted to 1:2000

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